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                 and Japanese-language basic patents from 2004-present
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NEWS 10 JAN 07 WPIDS, WPINDEX, and WPIX enhanced Japanese Patent
                 Classification Data
NEWS 11 FEB 02 Simultaneous left and right truncation (SLART) added
                 for CERAB, COMPUAB, ELCOM, and SOLIDSTATE
NEWS 12 FEB 02 GENBANK enhanced with SET PLURALS and SET SPELLING
NEWS 13 FEB 06 Patent sequence location (PSL) data added to USGENE
NEWS 14 FEB 10 COMPENDEX reloaded and enhanced
NEWS 15 FEB 11 WTEXTILES reloaded and enhanced
NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3,
             AND CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.
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STRUCTURE FILE UPDATES: 11 FEB 2009 HIGHEST RN 1104680-36-5 DICTIONARY FILE UPDATES: 11 FEB 2009 HIGHEST RN 1104680-36-5

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http://www.cas.org/support/stngen/stndoc/properties.html

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=> s gabapentin/cn
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T.1 1 GABAPENTIN/CN

=> d L1

ANSWER 1 OF 1 REGISTRY COPYRIGHT 2009 ACS on STN L1

RN 60142-96-3 REGISTRY Entered STN: 16 Nov 1984

ED

CN Cyclohexaneacetic acid, 1-(aminomethyl)- (CA INDEX NAME)

OTHER NAMES:

CN 1-(Aminomethyl)cyclohexaneacetic acid

CN CI 945

CN Gabapen

CN Gabapentin

CN Go 3450

CN GOE 2450

CN GOE 3450

CN Neurontin

ME C9 H17 N O2

CT COM

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(\*File contains numerically searchable property data) Other Sources: EINECS\*\*, WHO

(\*\*Enter CHEMLIST File for up-to-date regulatory information)

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT \*\*

2083 REFERENCES IN FILE CA (1907 TO DATE)
60 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
2096 REFERENCES IN FILE CAPLUS (1907 TO DATE)

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> d 12 20-23 IBIB ABS

L2 ANSWER 20 OF 23 CAPLUS COPYRIGHT 2009 ACS on STN ACCESSION NUMBER: 2004:331964 CAPLUS DOCUMENT NUMBER: 140:344917 TITLE: Gabapentin tablets preparation

INVENTOR(S): Manikandan, Ramalingam; Gogia, Ashish; Roy, Sunilendu

Bhushan; Malik, Rajiv

PATENT ASSIGNEE(S): Ranbaxy Laboratories Limited, India

SOURCE: PCT Int. Appl., 18 pp. CODEN: PIXXD2

DOCUMENT TYPE: Pat.ent.

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PA	TENT	NO.			KIND DATE					APPL	ICAT	DATE								
WO	2004	A1 20040422				WO 2	003-	IB44		2	0031	008								
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		GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	KZ,	LC,	LK,			
		LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NI,	NO,	NZ,			
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		TN,	TR,	TT,	TZ,	UA,	UG,	US,	UΖ,	VC,	VN,	YU,	ZA,	ZM,	zw					
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		FI,	FR,	GB,	GR,	ΗU,	ΙE,	IT,	LU,	MC,	NL,	PT,	RO,	SE,	SI,	SK,	TR,			
		BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	ΝE,	SN,	TD,	TG			
IN	IN 195214						2005	0128	IN 2002-DE1023						20021008					
	2002																			
	AU 2003267732					A1 20040504									CA, CH, CN, GE, GB, GD, GE, GB, GD, GE, KZ, LC, LK, NI, NO, NZ, SY, TJ, TM, ZW, AM, AZ, BY, DK, EE, ES, SM, TD, TG 20021008  20031008 20031008 20031008 S, MC, PT, HU, SK 20031008 20050407 20021008					
EP	1558	A1 20050803				EP 2	003-	7484		2	20031008 CA, CH, CN, GB, GD, GE, KZ, LC, LK, NI, NO, NZ, SY, TJ, TM, ZM, AZ, BY, DK, EE, ES, SI, SK, TR, SN, TD, TG 20031008 20031008 20031008 20031008 20031008 20031008 20050407 20021008									
	R:																PT,			
										CN 2003-80105033										
US	2006	0039	968		A1	A1 20060223				US 2005-530592						20050407				
PRIORIT	Y APP	LN.	INFO	.:						IN 2002-DE1023										
										WO 2	003-	IB44		W 20031008						

The present invention is generally directed to methods for preparing stable AB gabapentin tablets by wet granulation. A wet granulation method for preparing gabapentin tablets includes forming a mixture by dry mixing of a first portion of a binder with the gabapentin, one or more excipients, or a combination of the gabapentin and the one or more excipients; and adding a second portion of the binder to the mixture, wherein the second portion of the binder is in the form of a solution or

dispersion. REFERENCE COUNT:

THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD, ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 21 OF 23 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2003:319273 CAPLUS

DOCUMENT NUMBER: 138:326578

TITLE: Process for preparing tannate tablet, capsule or other

solid dosage forms INVENTOR(S): Kiel, Jeffrey S.; Thomas, H. Greg; Mani, Narasimhan

Kiel Laboratories, Inc., USA U.S. Pat. Appl. Publ., 7 pp. PATENT ASSIGNEE(S):

SOURCE:

CODEN: USXXCO

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 4

PATENT INFORMATION:

KIND DATE APPLICATION NO. DATE PATENT NO. US 20030077321 A1 20030424 US 2002-269027 20021010 US 7273623 B2 20070925

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CA 2453256 A1 20031023 CA 2003-2453256 20030226
CA 2469736 A1 20031023 CA 2003-2469736 20030226
W0 2003086356 A1 20031023 W0 2003-US5664 20030226
W: AU, CA, US
            WO 2003086346 A1 20031023 WO 2003-US5667
                                                                                                                                                         20030226
                   W: AU, CA, US
           AU 2003217703 AI 20031027 AU 2003-217703
AU 2003217704 AI 20031027 AU 2003-217704
AC 2482013 AI 20040422 CA 2003-02482013
WO 2004032826 A2 20040422 WO 2003-US10918
WO 2004032826 A3 20040826
                                                                                                                                                         20030226
                                                                                                                                                          20030226
                                                                                                                                                         20030409
                                                                                                                                                          20030409
                    W: CA
                    RW: AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
                                        A2 20060208 EP 2003-817708 20030409
            EP 1622586
                    R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
                              IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, SK
                                                                                                   US 2006-501649 20060809 US 2001-328990P P 20011012 US 2001-2828969P P 20011012 US 2002-282958 A 20020409 W 2003-US5664 W 20030226 WO 2003-US5667 W 2003-US10010 P 2003-US10
US 20050069584 A1 20050331 US 2004-503347 US 20070020332 A1 20070125 US 2006-501649 PRIORITY APPLN. INFO: US 2001-328990P
                                                                                                     WO 2003-US10918 W 20030409
US 2004-921438 A2 20040819
AB An active pharmaceutical ingredient is combined with tannic acid to form a
            tannate salt complex of the active ingredient. The active ingredient
           tannate salt complex without isolation or purification is then blended with
            pharmaceutically acceptable excipients to form a granulate which
            is processed into a tablet or capsule to generate a therapeutic solid
           dosage form. For example, tablets were prepared containing carbetapentane
            tannate 60.0 mg, chlorpheniramine tannate 4.0 mg, phenylephrine tannate
           10.0 mg, magnesium aluminum silicate 30.0 mg, Avicel PH 102 459.642 mg,
           Methocel E-10 M 5.0 mg, corn starch 3.0 mg, calcium phosphate dibasic
           10.133 mg, xanthan gum 7.875 mg, talc 2.25 mg, FD&C Red #40 0.85 mg, and
           magnesium stearate 2.25 mg.
REFERENCE COUNT:
                                                    33
                                                                        THERE ARE 33 CITED REFERENCES AVAILABLE FOR THIS
                                                                       RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L2 ANSWER 22 OF 23 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2002:256071 CAPLUS
DOCUMENT NUMBER:
                                                       136:284459
TITLE: STADLE SOLID GOODS:
INVENTOR(S): Spireas, Spiridon
PATENT ASSIGNEE(S): Sigmapharm, Inc., USA
SOURCE: PCT Int. Appl., 44 pp.
TITLE:
                                                       Stable solid dosage forms of amino acids
                                                        CODEN: PIXXD2
DOCUMENT TYPE: Patent LANGUAGE: English
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	PAT	ENT	NO.			KIN	D	DATE			APPL	ICAT:	DATE									
							_															
WO 2002026263						A2		20020404			WO 2	001-	20010926									
	WO 2002026263					A3		2003	0030103													
		W:	ΑE,	AG,	AL,	AM,	ΑT,	AU,	AZ,	BA,	BB,	BG,	BR,	BY,	BZ,	CA,	CH,	CN,				
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			GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	KZ,	LC,	LK,	LR,				
			LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	NZ,	PH,	PL,				
			PT.	RO.	RU.	SD.	SE.	SG.	ST.	SK.	SL.	T.T.	TM.	TR.	TT.	TZ.	IIA.	HG.				

FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION:

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                DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
                BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
      US 20020091159 A1 20020711 US 2001-928467
      US 7056951
                              B2 20060606
      CA 2422871 A1 20020404 CA 2001-2422871 20010926
AU 2001094736 A 20020408 AU 2001-94736 20010926
EP 1322335 A2 20030702 EP 2001-975405 20010926
           R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
                IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
NZ 524463 A 20041029 NZ 2001-524463
AU 2001294736 B2 20060817 AU 2001-294736
CN 1287782 C 20061206 CN 2001-816136
MX 2003002533 A 20040910 MX 2003-2553
PRIORITY APPLN. INFO:: U5 2000-235349P
                                     20041029 NZ 2001-524463
                                                                                   20010926
                                                      CN 2001-816136 20010926
MX 2003-2553 20030325
US 2000-235349P P 20000926
US 2001-928467 A 20010813
WO 2001-US30095 W 20010926
OTHER SOURCE(S):
                             MARPAT 136:284459
AB Pharmaceutical formulations contain an amino acid which is susceptible to
      the formation of an undesirable lactam, and a stabilizer comprising a
      volatile alc., a nonvolatile alc., a water-immiscible liquid or solid, a
      liquid with a relatively low dielec, constant, liquid and solid surfactants, an
      antioxidant, a ketone, an aldehyde, a solid polyethylene glycol of high
      mol. weight, polyvinylpyrrolidone, a derived cellulose, silicon dioxide, or a
      combination to inhibit the lactam formation. Thus, a formulation
      contained anhydrous gabapentin 400, corn starch 113, and water 100 mg/unit
      dose.
REFERENCE COUNT:
                                       THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS
                                       RECORD, ALL CITATIONS AVAILABLE IN THE RE FORMAT
L2 ANSWER 23 OF 23 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1999:753060 CAPLUS
DOCUMENT NUMBER:
                              131:356133
TITLE:
                              Solid compositions containing \( \gamma - \text{aminobutyric} \)
                              acid derivatives
INVENTOR(S):
                             Aomatsu, Akira
NOUNCE: Warner-Lambert Company, USA SOURCE: PCT Int. Appl., 99 pp.
                              CODEN: PIXXD2
                             Patent
DOCUMENT TYPE:
                             English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:
                        KIND DATE APPLICATION NO. DATE
      PATENT NO.
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      WO 9959572 A1 19991125 WO 1999-US10186 19990510
           W: AE, AL, AU, BA, BB, BG, BR, CA, CN, CU, CZ, EE, GD, GE, HR, HU,
                AB, AL, AU, BB, BB, BB, BR, CR, CL, CU, CL, EB, GU, MG, BL, HR, HO, 
ID, IL, IN, IS, JP, RF, RR, LC, LK, LR, II, LV, MG, MK, MM, MX, NN, NS, NO, NZ, FL, RO, SG, SI, SK, SL, TR, TT, UA, US, UZ, VN, YU, ZA, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
           RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK,
                ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, FT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
      CA 2325045
                               A1 19991125
                                                      CA 1999-2325045
                                                                                   19990510
     A1 19991125 CA 1999-2325045
CA 2325045 C 20050503
AU 9940733 A 19991206 AU 1999-40733
AU 769038 B2 20040115
BR 9910494 A 20010109 BR 1999-10494
EP 1077691 A1 20010228 EP 1999-924164
EP 1077691 B1 20080910
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		0110					A2 20011028 HU 2001-1791								19990510							
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		0000		1			A 20020415 EE 2000-671								19990510							
	507162					A			1128						62							
	1171587					С			1020						78							
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11	1 20	2001MN00451				A		2005	0318									20010424				
HF	( 10	3640	)7			A1		2005	0603	1	HK	200	1-3	1073	02		- 2	20011	018			
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JF	39	441	50			B2		2007	0711													
HF	( 10	7272	29			A1		2002	0418	1	HK	200	5-3	1054	87		- 2	20050	629			
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										1	OW	199	9-1	JS10	186		W :	19990	510			
											JP	200	2-:	1897	68		A3 :	19990	514			

## OTHER SOURCE(S): MARPAT 131:356133

AB The present invention provides a stabilized solid composition containing a 4-amino-3-substituted-butanoic acid derivative which can be obtained by incorporating a humectant as a stabilizer. Bulk powders of gabapentin (250 g) were sprayed with 72 g water by means of a fluidized granulator and then dried to give gabapentin granular powders A. A second batch of bulk powders of gabapentin (250 g) were sprayed with a solution of 5 g propylene glycol in 67 g water by means of the fluidized granulator and then dried to give gabapentin granular powders B. The gabapentin granular powders A and B obtained were stored under conditions and then the lactam formed in each of the powders was determined by HPLC. E.g., gabapentin bulk powders stored for 4 wk at 50° and 85% humidity did not show any degradation

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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